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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07D 409/04, A61K 31/445		A1	(11) International Publication Number: WO 96/16961
			(43) International Publication Date: 6 June 1996 (06.06.96)
(21) International Application Number: PCT/US95/15320 (22) International Filing Date: 22 November 1995 (22.11.95) (30) Priority Data: 08/345,188 28 November 1994 (28.11.94) US (71) Applicant: AMERICAN HOME PRODUCTS CORPORATION [US/US]; Five Giralda Farms, Madison, NJ 07940-0874 (US). (72) Inventor: BAUDY, Reinhardt, Baudy; 1483 Scarlet Oak Road, Yardley, PA 19067 (US). (74) Agents: ALICE, Ronald, W.; American Home Products Corporation, Five Giralda Farms, Madison, NJ 07940-0874 (US) et al.		(81) Designated States: AL, AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU, IS, JP, KG, KP, KR, KZ, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TT, UA, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, LS, MW, SD, SZ, UG). Published With international search report.	
(54) Title: (THIOPHEN-2-YL)-PIPERIDIN OR TETRAHYDROPYRIDIN CARBOXAMIDES			
(57) Abstract This invention provides compounds having formula (I), wherein R is hydrogen, alkyl, alkenyl, alkynyl, -COR ² , phenyl, or phenylalkyl of 7-10 carbon atoms; the dotted line represents an optional double bond; R ¹ is hydrogen, -OH, OR ³ , or is absent if the optional double bond is present; R ² and R ³ are each, independently, alkyl, alkenyl, alkynyl, phenyl, or phenylalkyl; R ⁴ is hydrogen, -OR ⁵ , alkyl, alkenyl, alkynyl, -COR ⁵ , -CO ₂ R ⁵ , -CONR ⁵ R ⁶ , perhaloalkyl, halogen, phenyl, or phenylalkyl; R ⁵ and R ⁶ are each, independently, hydrogen, alkyl, alkenyl, alkynyl, phenyl, or phenylalkyl; and n = 0 - 2 or a pharmaceutically acceptable salt thereof that are useful as antipsychotic, antidepressant and anxiolytic agents useful in the treatment and relief of the symptoms of these disease states.			
<div style="text-align: right;">(I)</div>			

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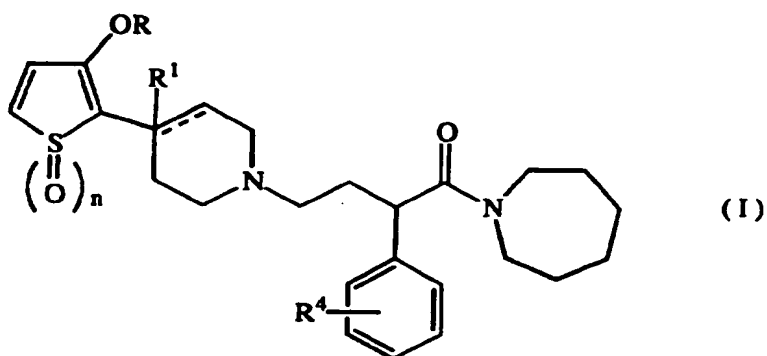
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(THIOPHEN-2-YL)-PIPERIDIN OR TETRAHYDROPYRIDIN
CARBOXAMIDES

- 5 This invention provides compounds having selectivity for the serotonergic 5-HT_{1A} receptor, useful in the treatment of central nervous system disorders, having the formula I



- 10 wherein
 R is hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR², phenyl, or phenylalkyl of 7-10 carbon atoms;
 the dotted line represents an optional double bond;
 R¹ is hydrogen, -OH, OR³, or is absent if the optional double bond is present;
 15 R² and R³ are each, independently, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms;
 R⁴ is hydrogen, -OR⁵, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR⁵, -CO₂R⁵, -CONR⁵R⁶, perhaloalkyl of 1-6 carbon atoms, halogen, phenyl, or phenylalkyl of 7-10 carbon atoms;
 20 R⁵ and R⁶ are each, independently, hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms; and
 n = 0 - 2
 25 or a pharmaceutically acceptable salt thereof.

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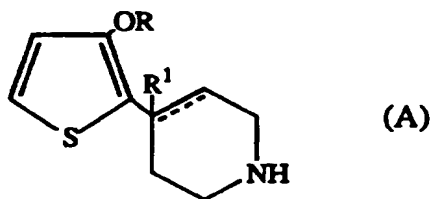
The pharmaceutically acceptable salts are those derived from such organic and inorganic acids as: acetic, lactic, citric, tartaric, succinic, maleic, malonic, gluconic, hydrochloric, hydrobromic, phosphoric, nitric, sulfuric, methanesulfonic, and similarly known acceptable acids.

5 The terms alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, and alkynyl of 2-7 carbon atoms, include both straight chain as well as branched carbon chains. In the generic structure described above, when $n = 0$, the sulfur containing ring is a thiophene ring, when $n = 1$, the sulfur containing ring is a thiophene S-oxide, and when $n = 2$, the sulfur containing ring is a thiophene S-dioxide. The term "halogen" refers to fluoro, chloro, bromo, or iodo.

10 The compounds within the scope of the invention by virtue of their configuration, exhibit stereoisomerism. Such centers can contain either the R or S configuration or can be racemic with respect to such center or centers. Accordingly, the compounds of the invention include the diastereomers, enantiomers, racemates and mixtures thereof.

Of these compounds, the preferred members are those in which $n = 0$; and those in which $n = 0$, and R is alkyl of 1-6 carbon atoms.

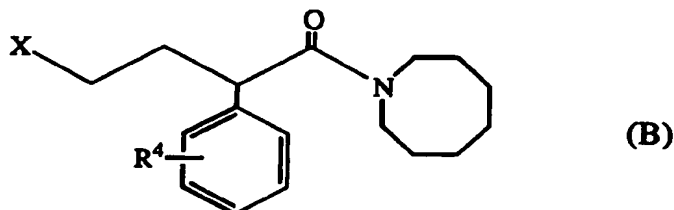
20 The compounds of this invention can be prepared by conventional methods. For example, the compounds where n is 1 or 2 may be made by oxidation of compounds where n is 0. The compounds where n is 0 may be prepared by a process which comprises subjecting a compound having the formula A



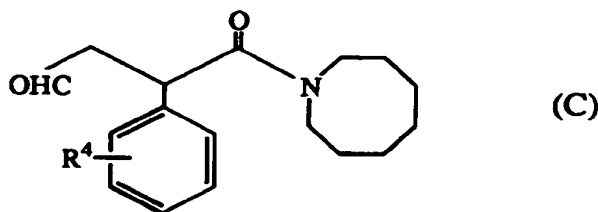
25 where R, R^1 and the dotted line are as defined above to

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(a) alkylation by reaction with a compound having the formula B



where R⁴ is as defined above and X is a leaving group, preferably chloro or bromo, or
 (b) reductive alkylation with an aldehyde having the formula C



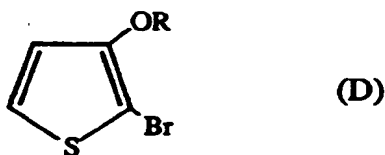
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where R⁴ is as defined above; and, where desired, a compound with one meaning of R¹ may be converted into a compound with another meaning of R¹ as an after process.

The alkylation process (a) is preferably carried out in the presence of a base, for example diisopropyl ethylamine. The reductive alkylation of the aldehyde (b) may be carried out with sodium borohydride as reducing agent.

10

The compounds having the formula A may be prepared by a process which includes the step of reacting a compound having the formula D



15

where R is as defined above or a protecting group is subjected to a Grignard reaction with addition of an N-protected 4-piperidone to afford an N-protected-4-hydroxy-4-(3-OR-thiophen-2-yl)piperidine. The N-protecting group may be in the form of a carbamate, preferably ethyl carbamate. The free piperidine nitrogen atom may be obtained by removal of the protecting group, for example, using KOH in the case of a carbamate. Where -OR represents a protected hydroxy group, the protection may be

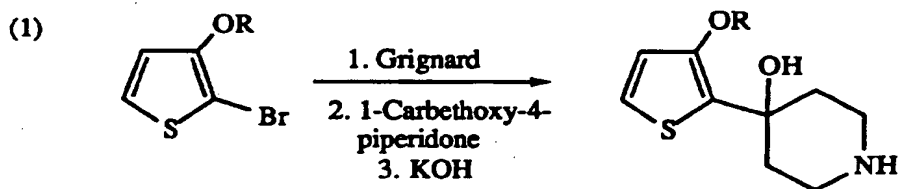
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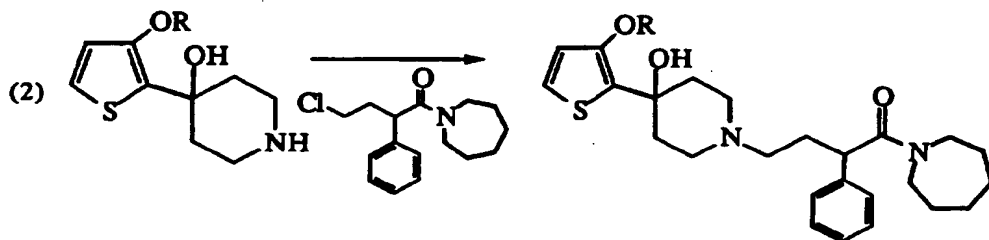
removed in an after-step. Again a protecting group which can be removed under basic (or neutral) conditions is preferred. This is because the tertiary alcohol is liable to undergo dehydration under acidic conditions.

Before, during or after the alkylation step (a) or reductive alkylation step (b) one may convert one meaning of R^1 into another. For example a 4-hydroxypiperidine may be converted into a tetrahydropyridine derivative by dehydration. That derivative may be reduced where a compound in which R^1 is hydrogen is desired. Ethers may be prepared in a known manner.

As an example of a synthesis of compounds of the invention the appropriately substituted 2-bromo-thiophene can be subjected to a Grignard reaction in which the addition to a 4-piperidone carbamate affords the desired tertiary alcohol. Subsequent hydrolysis of the carbamate yields the desired 4-hydroxy-4-thiophen-2-yl-piperidine (1).



The 4-substituted piperidine can then be alkylated with a substituted 4-halo-butanoic acid amide in the presence of an acid scavenger such as diisopropylethylamine to yield the final product (2).



Further functionalization of the piperidine hydroxyl group can be accomplished using standard methodology, and dehydration of the hydroxyl group to provide tetrahydropyridine derivatives can be accomplished under mildly acidic conditions.

Compounds of this invention in which n is 1 or 2 can be prepared by oxidizing a compound of formula I in which n is 0 with an oxidizing agent such as *m*-chloroperbenzoic acid. When n is not equal to 0, it is preferred that n equal 1.

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Representative compounds of this invention were evaluated and determined to have high affinity for the serotonin 5-HT_{1A} receptor by evaluating the compound's ability to displace [³H] 8-OHDPAT (dipropylaminotetralin) from the 5-HT_{1A} serotonin receptor following the procedure of Hall et al., J. Neurochem. 44, 1685 (1985). This standard pharmacological test procedure was employed to analogize this property of the claimed compounds with that of buspirone, which is a standard for anxiolytic activity, and, like the compounds of this invention, displays potent affinity for the 5-HT_{1A} serotonin receptor subtype. The anxiolytic activity of buspirone is believed to be, at least partially, due to its 5-HT_{1A} receptor affinity (Vander Maclen et al., Eur. J. Pharmacol. 1986, 129 (1-2) 133-130). In this standard pharmacological test procedure, buspirone has an IC₅₀ of approximately 10 nM.

The results obtained for representative compounds of this invention in the standard pharmacological test procedure described above, are as follows:

Compound	5-HT _{1A} Binding (IC ₅₀)
Example 1	0.8 nM
Example 2	0.25 nM
Example 3	0.98 nM
Example 4	0.2 nM
Example 5	0.7 nM

The results obtained in the standard pharmacological test procedure demonstrate that the compounds this invention possess high affinities for the serotonin 5-HT_{1A} receptor, and consequently, they are useful in the treatment of multi-CNS disorders amenable to treatment with antipsychotic, antidepressant and anxiolytic agents. As such, the compounds of this invention may be administered a mammal in need of antipsychotic, antidepressant and/or anxiolytic medical treatment in an amount sufficient to alleviate the symptoms of the disease state, such as depression, paranoia, schizophrenia, anxiety, sleep disorders, eating disorders, cognitive disorders, panic, social phobia, obsessive compulsive disorders, sexual dysfunction, addiction, and related problems. When administered for the treatment of the above disease states, the compounds of this invention can be administered to a mammal orally, parenterally, intranasally, intrabronchially, transdermally, intravaginally, or rectally.

35

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The compounds of this invention can be formulated neat or with a pharmaceutical carrier to a mammal in need thereof. The pharmaceutical carrier may be solid or liquid.

5 A solid carrier can include one or more substances which may also act as flavoring agents, lubricants, solubilizers, suspending agents, fillers, glidants, compression aids, binders or tablet-disintegrating agents; it can also be an encapsulating material. In powders, the carrier is a finely divided solid which is in admixture with the finely divided active ingredient. In tablets, the active ingredient is mixed with a carrier having the necessary compression properties in suitable proportions and compacted in
10 the shape and size desired. The powders and tablets preferably contain up to 99% of the active ingredient. Suitable solid carriers include, for example, calcium phosphate, magnesium stearate, talc, sugars, lactose, dextrin, starch, gelatin, cellulose, methyl cellulose, sodium carboxymethyl cellulose, polyvinylpyrrolidone, low melting waxes and ion exchange resins.

15 Liquid carriers are used in preparing solutions, suspensions, emulsions, syrups, elixirs and pressurized compositions. The active ingredient can be dissolved or suspended in a pharmaceutically acceptable liquid carrier such as water, an organic solvent, a mixture of both or pharmaceutically acceptable oils or fats. The liquid carrier can contain other suitable pharmaceutical additives such as solubilizers, emulsifiers,
20 buffers, preservatives, sweeteners, flavoring agents, suspending agents, thickening agents, colors, viscosity regulators, stabilizers or osmo-regulators. Suitable examples of liquid carriers for oral and parenteral administration include water (partially containing additives as above, e.g. cellulose derivatives, preferably sodium carboxymethyl cellulose solution), alcohols (including monohydric alcohols and
25 polyhydric alcohols, e.g. glycols) and their derivatives, and oils (e.g. fractionated coconut oil and arachis oil). For parenteral administration, the carrier can also be an oily ester such as ethyl oleate and isopropyl myristate. Sterile liquid carriers are useful in sterile liquid form compositions for parenteral administration. The liquid carrier for pressurized compositions can be halogenated hydrocarbon or other pharmaceutically
30 acceptable propellant.

Liquid pharmaceutical compositions which are sterile solutions or suspensions can be utilized by, for example, intramuscular, intraperitoneal or subcutaneous injection. Sterile solutions can also be administered intravenously. The compound can also be administered orally either in liquid or solid composition form.

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The compounds of this invention may be administered rectally in the form of a conventional suppository. For administration by intranasal or intrabronchial inhalation or insufflation, the compounds of this invention may be formulated into an aqueous or partially aqueous solution, which can then be utilized in the form of an aerosol. The compounds of this invention may also be administered transdermally through the use of a transdermal patch containing the active compound and a carrier that is inert to the active compound, is non toxic to the skin, and allows delivery of the agent for systemic absorption into the blood stream via the skin. The carrier may take any number of forms such as creams and ointments, pastes, gels, and occlusive devices. The creams and ointments may be viscous liquid or semisolid emulsions of either the oil-in-water or water-in-oil type. Pastes comprised of absorptive powders dispersed in petroleum or hydrophilic petroleum containing the active ingredient may also be suitable. A variety of occlusive devices may be used to release the active ingredient into the blood stream such as a semipermeable membrane covering a reservoir containing the active ingredient with or without a carrier, or a matrix containing the active ingredient. Other occlusive devices are known in the literature.

The dosage to be used in the treatment of a specific psychosis must be subjectively determined by the attending physician. The variables involved include the specific psychosis or state of anxiety or depression and the size, age and response pattern of the patient. Based on the results obtained in the standard pharmacological test procedures, projected oral daily dosages of active compound would be 1-100 mg/kg, preferably between 1-30 mg/kg, and more preferably between 1-10 mg/kg. Projected intravenous daily dosages would be 0.2-20 mg/kg, preferably between 0.2-6 mg/kg, and more preferably between 0.2-2 mg/kg. Treatment will generally be initiated with small dosages less than the optimum dose of the compound. Thereafter the dosage is increased until the optimum effect under the circumstances is reached; precise dosages for oral, parenteral, nasal, or intrabronchial administration will be determined by the administering physician based on experience with the individual subject treated. Preferably, the pharmaceutical composition is in unit dosage form, e.g. as tablets or capsules. In such form, the composition is sub-divided in unit dose containing appropriate quantities of the active ingredient; the unit dosage forms can be packaged compositions, for example, packeted powders, vials, ampoules, prefilled syringes or sachets containing liquids. The unit dosage form can be, for example, a capsule or tablet itself, or it can be the appropriate number of any such compositions in package form.

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The following examples illustrate the production of representative compounds of this invention.

5

Example 1**1-Azepan-1-yl-4-[4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one 0.75 hydroiodide.**

A mixture of 4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidine (2.4 g, 11 mmole), 1-(azepan-1-yl)-4-chloro-2-phenyl-butan-1-one (3.63 g, 13 mmole),
10 diisopropyl-ethylamine (1.68 g, 13 mmole) and potassium iodide (2.16 g, 13 mmole) was heated in dimethylformamide (40 mL) to 80°C for 4 hours. After cooling to ambient temperature the solvent was evaporated in vacuo and the residue suspended in ethyl acetate (80 mL). The precipitate was filtered and recrystallized from acetonitrile to yield 4.3 g of product, m.p. 126-8 °C.

15

Elemental Analysis for: C₂₆H₃₆N₂O₃S • 0.75 HI.

Calcd: C, 56.51; H, 6.70; N, 5.07.

Found: C, 56.28; H, 6.43; N, 4.88.

20

Example 2**1-Azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one**

The title compound was prepared from 1-azepan-1-yl-4-[4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one (1.35 g, 2.9 mmole) and
25 acetic acid (80 mL) in the manner previously described above to yield 1.1 g of product, m.p. 108-12 °C.

Elemental Analysis for: C₂₆H₃₄N₂O₂S • 0.1 chloroform.

Calcd: C, 69.49; H, 7.64; N, 6.23.

30 **Found:** C, 69.83; H, 7.47; N, 6.10.

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Example 3

1-Azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-
piperidin-1-yl]-2-phenyl-butan-1-one fumarate salt

The title compound was prepared from 1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one (1 g, 2.2 mmole), acetic acid (50 mL), and Pearlman's catalyst (0.3 g) in the manner previously described above to yield 0.6 g of product, m.p. 134-8 °C. The title compound was isolated as a hemihydrate.

10 Elemental Analysis for: $C_{26}H_{36}N_2O_2S \cdot C_8H_8O_8 \cdot 0.5 H_2O$.

Calcd: C, 59.90; H, 6.65; N, 4.11.

Found: C, 59.58; H, 6.71; N, 4.11.

Example 4

15 (S)-1-Azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-
tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one hydrochloride salt

Acetic acid (0.48 mL) was added to a solution of the starting 4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidine (853 mg, 4 mmole) and (S)-4-(azepan-1-yl)-4-oxo-3-phenyl-butyraldehyde (1037 mg, 4 mmole) in methanol (20 mL). At ambient temperature the reaction mixture was treated portionwise with sodium cyanoborohydride (276 mg, 4.4 mmole). The mixture was stirred an additional three hours at 25°C and then poured into aqueous saturated sodium bicarbonate solution (50 mL). The aqueous phase was extracted with methylene chloride (3 x 50 mL). The combined organic extracts were washed with water (100 mL), brine (80 mL), dried over magnesium sulfate and filtered. The filtrate was evaporated in vacuo and the residue dissolved in chloroform (50 mL). Ethanolic hydrochloric acid was added and the resulting solution evaporated. The residue was triturated in ether and the precipitate filtered and dried to yield 400 mg of the title compound, mp 114-6°C. The title compound was isolated as the hydrate.

30

Elemental Analysis for: $C_{26}H_{34}N_2O_2S \cdot 1 HCl \cdot 1 H_2O$.

Calcd: C, 63.30; H, 7.56; N, 5.68.

Found: C, 63.20 H, 7.68; N, 5.66.

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Example 5**(2S)-1-(Azepan-1-yl)-4-[4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one dihydrochloride salt**

5 A solution of (S)-1-Azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one (600 mg, 1.3 mmole) in glacial acetic acid (100 mL) was treated at once with Pearlman's Catalyst (600 mg) and hydrogenated at 50 psi at ambient temperature for 12 hours. After purging the reaction vessel with nitrogen the catalyst was removed by filtration, washed with acetic acid (30 mL) and the filtrate evaporated in vacuo. The residue was partitioned between 5% aqueous
10 sodium bicarbonate (50 mL) and chloroform (100 mL). The separated organic layer was dried over magnesium sulfate, filtered and the filtrate evaporated to dryness in vacuo. Column chromatography on 20 g of silica gel with 3% methanol / chloroform as eluant, followed by trituration in ether with the addition of ethanolic hydrochloric acid gave 110 mg of the title compound, m.p. 91-3°C.

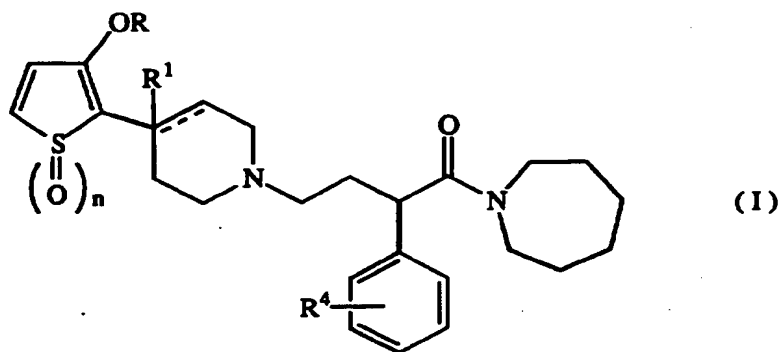
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Elemental Analysis for: $C_{26}H_{36}N_2O_2S \cdot 2 HCl$ **Calcd:** C, 60.80; H, 7.45; N, 5.45.**Found:** C, 61.19; H, 7.75; N, 5.43.

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What is claimed is

1. A compound having the formula (I)



5

wherein

- R is hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR², phenyl, or phenylalkyl of 7-10 carbon atoms;
- 10 the dotted line represents an optional double bond;
- R¹ is hydrogen, -OH, OR³, or is absent if the optional double bond is present;
- R² and R³ are each, independently, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms;
- 15 R⁴ is hydrogen, -OR⁵, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR⁵, -CO₂R⁵, -CONR⁵R⁶, perhaloalkyl of 1-6 carbon atoms, halogen, phenyl, or phenylalkyl of 7-10 carbon atoms;
- R⁵ and R⁶ are each, independently, hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms; and
- 20 n = 0 - 2
- or a pharmaceutically acceptable salt thereof.

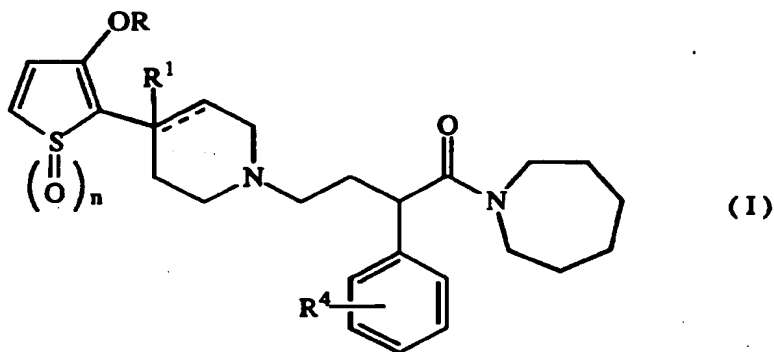
2. The compound of claim 1, wherein n = 0 or a pharmaceutically acceptable salt thereof.
- 25

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3. The compound of claim 2, wherein R is alkyl of 1-6 carbon atoms or a pharmaceutically acceptable salt thereof.
4. The compound of claim 1 which is 1-azepan-1-yl-4-[4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one or a pharmaceutically acceptable salt thereof.
- 5 5. The compound of claim 1 which is 1-azepan-1-yl-4-[4-hydroxy-4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one hydroiodide salt.
- 10 6. The compound of claim 1 which is 1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one or a pharmaceutically acceptable salt thereof.
- 15 7. The compound of claim 1 which is 1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one or a pharmaceutically acceptable salt thereof.
8. The compound of claim 1 which is 1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one fumarate salt.
- 20 9. The compound of claim 1 which is (S)-1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one or a pharmaceutically acceptable salt thereof.
- 25 10. The compound of claim 1 which is (S)-1-azepan-1-yl-4-[4-(3-methoxy-thiophen-2-yl)-1,2,3,6-tetrahydro-pyridin-1-yl]-2-phenyl-butan-1-one hydrochloride salt.
- 30 11. A compound of claim 1 which is (2S)-1-(azepan-1-yl)-4-[4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one or a pharmaceutically acceptable salt thereof.
- 35 12. A compound of claim 1 which is (2S)-1-(azepan-1-yl)-4-[4-(3-methoxy-thiophen-2-yl)-piperidin-1-yl]-2-phenyl-butan-1-one dihydrochloride salt.

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13. A method of treating anxiety, psychosis, or depression in a mammal in need thereof which comprises administering to said mammal, an effective amount of a compound of the formula (I)



5

wherein

R is hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR², phenyl, or phenylalkyl of 7-10 carbon atoms;

10 the dotted line represents an optional double bond;

R¹ is hydrogen, -OH, OR³, or is absent if the optional double bond is present;

R² and R³ are each, independently, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms;

15 R⁴ is hydrogen, -OR⁵, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, -COR⁵, -CO₂R⁵, -CONR⁵R⁶, perhaloalkyl of 1-6 carbon atoms, halogen, phenyl, or phenylalkyl of 7-10 carbon atoms;

R⁵ and R⁶ are each, independently, hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms; and

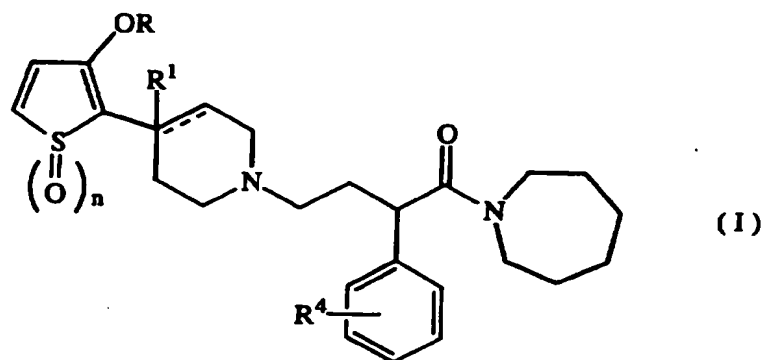
20

n = 0 - 2

or a pharmaceutically acceptable salt thereof.

- 14 -

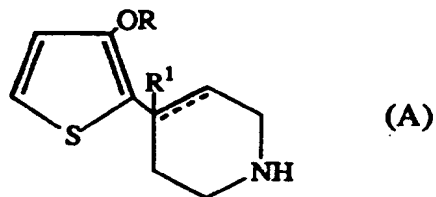
14. A pharmaceutical composition which comprises a compound of the formula (I)



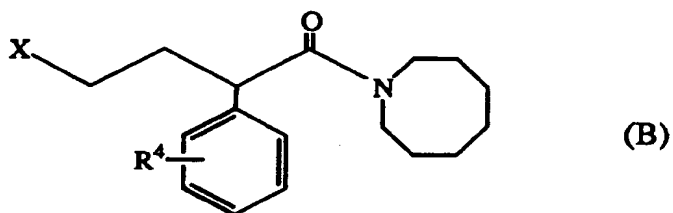
- 5 wherein
- R is hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, $-\text{COR}^2$, phenyl, or phenylalkyl of 7-10 carbon atoms; the dotted line represents an optional double bond;
- R^1 is hydrogen, $-\text{OH}$, OR^3 , or is absent if the optional double bond is present;
- 10 R^2 and R^3 are each, independently, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms;
- R^4 is hydrogen, $-\text{OR}^5$, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, $-\text{COR}^5$, $-\text{CO}_2\text{R}^5$, $-\text{CONR}^5\text{R}^6$, perhaloalkyl of 1-6 carbon atoms, halogen, phenyl, or phenylalkyl of 7-10 carbon atoms;
- 15 R^5 and R^6 are each, independently, hydrogen, alkyl of 1-6 carbon atoms, alkenyl of 2-7 carbon atoms, alkynyl of 2-7 carbon atoms, phenyl, or phenylalkyl of 7-10 carbon atoms; and
- $n = 0 - 2$
- 20 or a pharmaceutically acceptable salt thereof, and a pharmaceutical carrier.

- 15 -

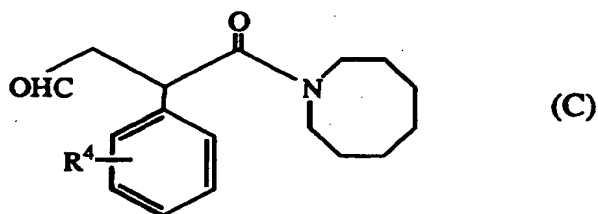
15. A process for preparing compounds of formula I, as claimed in Claim 1, where n is 0, by subjecting a compound having the formula A



5 where R, R¹ and the dotted line are as defined above to
 (a) alkylation by reaction with a compound having the formula B



where R⁴ is as defined above and X is a leaving group, preferably chloro or bromo, or
 (b) reductive alkylation with an aldehyde having the formula C



10

where R⁴ is as defined above; and, where desired, a compound with one meaning of R¹ may be converted into a compound with another meaning of R¹ as an after process.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 95/15320

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07D409/04 A61K31/445

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO,A,94 03444 (JOHN WYETH & BROTHER LTD.) 17 February 1994 see the whole document ---	1-15
A	EP,A,0 595 166 (BAYER AG.) 4 May 1994 see the whole document -----	1-15

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- * "A" document defining the general state of the art which is not considered to be of particular relevance
- * "E" earlier document but published on or after the international filing date
- * "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- * "O" document referring to an oral disclosure, use, exhibition or other means
- * "P" document published prior to the international filing date but later than the priority date claimed

- * "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- * "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- * "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- * "Z" document member of the same patent family

Date of the actual completion of the international search

2 February 1996

Date of mailing of the international search report

14.02.96

Name and mailing address of the ISA

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Authorized officer

Chouly, J

INTERNATIONAL SEARCH REPORT

In national application No.

PCT/US95/15320

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claim 13 is directed to a method of treatment of (diagnostic method practised on) the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 95/15320

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO-A-9403444	17-02-94	AU-B- 4714593	03-03-94
		CA-A- 2141810	17-02-94
		EP-A- 0664801	02-08-95
		FI-A- 950486	03-02-95

EP-A-595166	04-05-94	DE-A- 4236707	05-05-94
		AU-B- 4873093	12-05-94
		CA-A- 2109352	01-05-94
		CN-A- 1094403	02-11-94
		CZ-A- 9302276	18-05-94
		FI-A- 934778	01-05-94
		HU-A- 68941	28-08-95
		JP-A- 6220047	09-08-94
		NO-A- 933718	02-05-94
		SK-A- 120793	09-11-94
		ZA-A- 9308088	07-06-94
